

REPORTING AND PROCESSING MEDICAL MATERIEL COMPLAINTS/
QUALITY IMPROVEMENT REPORT

DATE

NO.

TO

FROM

TYPE OF COMPLAINT ►

1A. FOR DOD USE

☐ I☐ II☐ III

1B. FOR VA USE

☐ QUALITY COMPLAINT☐ NEW ITEM☐ SIMILAR ITEM

2. NATIONAL STOCK NO.

3. ITEM DESCRIPTION

4. NAME AND ADDRESS OF MANUFACTURER

5. NAME OF CONTRACTOR (If other than the manufacturer)

6. CONTRACT NO. OR PURCHASE ORDER NO.

7A. VA DEPOT VOUCHER NO.

7B. DOD REQUISITION NO.

8. LOT NO.

9. CONTROL NO.

10. MANUFACTURER'S SERIAL NO.

11. MODEL NO.

12. DATE MANUFACTURED

13. DATE PACKED

14. EXPIRATION DATE

15. SOURCE (Name of Depot)

16. QUANTITY ON HAND

17. QUANTITY SUSPENDED

COMPLETE ITEM 18A THROUGH F18F FOR DOD TYPE 1 COMPLAINTS ONLY

18A. TOTAL NO. PATIENTS INVOLVED

18B. TOTAL NO. REACTIONS

18C. SEVERE OR UNUSUAL REACTIONS

18D. REACTIONS REQUIRING
HOSPITALIZATION

18E. LENGTH OF HOSPITALIZATION

18F. VACCINE

☐ INITIAL☐ BOOSTER

INTERVAL _____

19. CAUSE OF COMPLAINT (Explanation of unsatisfactory condition, deficiency, or description of reaction. Complete 19 through 22 for ALL complaints.)

20A. TYPED NAME OF INITIATOR (For Type I MC/DC/NC)

20B. AUTOVON/FTS TELEPHONE NO.

20C. COMMERCIAL TELEPHONE NO.

21A. TYPED NAME OF SUPPLY OFFICER

21B. SIGNATURE OF SUPPLY OFFICER

21C. DATE

21D. AUTOVON/FTS TELEPHONE NO.

21E. COMMERCIAL TELEPHONE NO.

AREA CODE ()

REPORTING AND PROCESSING MEDICAL MATERIEL COMPLAINTS/QUALITY IMPROVEMENT REPORT (Continued)

22. RECOMMENDATIONS AND/OR ADDITIONAL REMARKS

23. ACTION TAKEN

24. NAME (*Action Officer*)

25. TITLE AND ORGANIZATION

26. DATE